IN THE CLAIMS:

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Please amend the claims in the subject patent application as follows:

- 1. (original) A syringe which is comprised of a barrel having a fluid chamber, a proximal end, a distal end and an elongated tip extending from said distal end having a passageway therethrough in fluid communication with said chamber, and an elongated plunger rod including a stopper slidably positioned in fluid-tight engagement with an inside surface of said chamber for drawing fluid into and out of said chamber by movement of said plunger relative to said barrel, wherein the stopper is comprised of neodymium polyisoprene rubber.
 - 2. (original) A syringe as specified in claim 1 wherein the stopper has ribs.
- 3. (original) A syringe wherein the neodymium polyisoprene rubber is made with a catalyst system that includes (1) an organoaluminum compound, (2) an organoneodymium compound, and (3) at least one compound that contains at least one labile halide ion.
- 4. (currently amended) A syringe as specification in claim 3 which is comprised of a barrel having a fluid chamber, a proximal end, a distal end and an elongated tip extending from said distal end having a passageway therethrough in fluid communication with said chamber, and an elongated plunger rod including a stopper slidably positioned in fluid-tight engagement with an inside surface of said chamber for drawing fluid into and out of said chamber by movement of said plunger relative to said barrel, wherein the stopper is comprised of neodymium polyisoprene rubber, wherein the stopper has ribs, wherein the neodymium polyisoprene rubber is made with a catalyst system that includes (1) an organoaluminum compound, (2) an organoneodymium compound, and (3) at least one compound that contains at least one labile halide ion, and wherein the organoaluminum compound is of the structural formula:

wherein R_1 is selected from the group consisting of alkyl, alkoxy, aryl, alkaryl, arylalkyl radicals and hydrogen, wherein R_2 is selected from the group consisting of alkyl, aryl, alkaryl, arylalkyl radicals and hydrogen, and wherein R_3 is selected from a group consisting of alkyl, aryl, alkaryl and arylalkyl radicals.

- 5. (original) A syringe as specified in claim 3 wherein the organoaluminum compound is selected from the group consisting of trimethylaluminum, triethylaluminum, trin-n-propylaluminum, triisopropylaluminum, triisopropyla
- 6. (original) A syringe as specified in claim 3 wherein the organoneodymium compound is of the formula NdL₃ wherein Nd represents neodymium and L is an organic ligand selected from a group consisting of (1) o-hydroxyaldehydes, (2) o-hydroxyphenones, (3) aminophenols, (4) hydroxy esters, (5) hydroxy quinolines, (6) beta-diketones, (7) monocarboxylic acids, (8) ortho dihydric phenols, (9) alkylene glycols, (10) dicarboxylic acids, (11) alkylated derivatives of dicarboxylic acids, and (12) phenolic ethers.
- 7. (original) A syringe as specified in claim 3 wherein the organoneodymium compound is of the formula NdL₃ wherein L represents an organic ligand containing from 1 to 20 carbon atoms.
- 8. (original) A syringe as specified in claim 3 wherein the organoneodymium is neodymium naphthenate.
- 9. (original) A syringe as specified in claim 3 wherein the organoneodymium is neodymium neodecanoate.

- 10. (original) A syringe as specified in claim 3 wherein the organoneodymium is neodymium octanoate.
- 11. (original) A syringe as specified in claim 3 wherein the compound that contains at least one labile halide ion is selected from the group consisting of inorganic halide acids, organometallic halides, and inorganic halides.
- 12. (original) A syringe as specified in claim 11 wherein the compound that contains at least one labile halide ion is an inorganic halide acid selected from the group consisting of hydrogen bromide, hydrogen chloride, and hydrogen iodide.
- 13. (original) A syringe as specified in claim 11 wherein the compound that contains at least one labile halide ion is an organometallic halide selected from the group consisting of ethylmagnesium bromide, butylmagnesium bromide, phenylmagnesium bromide, methylmagnesium chloride, butylmagnesium chloride, ethylmagnesium iodide, phenylmagnesium iodide, diethylaluminum bromide, diisobutylaluminum bromide, methylaluminum sesquibromide, diethylaluminum chloride, ethylaluminum dichloride, ethylaluminum sesquichloride, diisobutylaluminum chloride, isobutylaluminum dichloride, dihexylaluminum chloride, cyclohexylaluminum dichloride, phenylaluminum dichloride, dioddecylaluminum chloride, diethylaluminum fluoride, dibutylaluminum fluoride, diethylaluminum iodide, dibutylaluminum iodide, phenylaluminum diiodide, trimethyltin bromide, triethyltin chloride, dibutyltin dichloride, butyltin trichloride, diphenyltin dichloride, and tributyltin iodide.
- 14. (original) A syringe as specified in claim 11 wherein the compound that contains at least one labile halide ion is an inorganic halide selected from the group consisting of aluminum bromide, aluminum chloride, aluminum iodide, antimony pentachloride, antimony trichloride, boron tribromide, boron trichloride, ferric chloride, gallium trichloride, molybdenum pentachloride, phosphorus tribromide, phosphorus pentachloride, stannic chloride, titanium tetrachloride, titanium tetrachloride, titanium tetrachloride.

- 15. (original) A syringe as specified in claim 3 wherein the atomic ratio of the halide ion to the neodymium is within the range of 0.1:1 to 6:1, and wherein the molar ratio of the organoaluminum compound to the organoneodymium compound is within the range of 4:1 to 200:1.
- 16. A syringe mixing and delivery system comprising a first barrel (original) having an open end and an opposite delivery end defining a delivery passage; a reciprocable stopper sealingly disposed in said first barrel to define a first chamber between said delivery passage and said reciprocable stopper for containing a first constituent in said first chamber; a second barrel that is sized to be disposed in said first barrel and that has an open end and an opposite discharge end defining a discharge passage; a slidable plunger sealingly disposed within said second barrel to define a second chamber between said discharge passage and said slidable plunger for containing a liquid second constituent in said second chamber; and fluid transfer connector means for operatively connecting said second barrel with said reciprocable stopper to permit flow of said liquid second constituent through said stopper from said second chamber to said first chamber to mix with said first constituent when said second barrel discharge end and plunger are moved closer together whereby subsequent movement of said second barrel and reciprocable stopper together toward said delivery passage of said first barrel expresses the mixed constituents out of said first chamber through said delivery passage, wherein said reciprocable stopper and plunger are comprised of neodymium polyisoprene rubber.
- 17. (original) A syringe mixing and delivery system as specified in claim 16, wherein said stopper has ribs.
- 18. (original) A syringe mixing and delivery system as specified in claim 17, wherein the neodymium polyisoprene rubber is made with a catalyst system that includes (1) an organoaluminum compound, (2) an organoneodymium compound, and (3) at least one compound that contains at least one labile halide ion.
 - 19. (new) A syringe as specification in claim 18 wherein the organoaluminum

compound is of the structural formula:

$$R_1-AI$$
 R_2

wherein R1 is selected from the group consisting of alkyl, alkoxy, aryl, alkaryl, arylalkyl radicals and hydrogen, wherein R2 is selected from the group consisting of alkyl, aryl, alkaryl, arylalkyl radicals and hydrogen, and wherein R3 is selected from a group consisting of alkyl, aryl, alkaryl and arylalkyl radicals.